

This Page Is Inserted by IFW Operations
and is not a part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

**As rescanning documents *will not* correct images,
please do not report the images to the
Image Problem Mailbox.**

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11)

EP 1 054 036 A1

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
22.11.2000 Bulletin 2000/47

(51) Int Cl.7: **C08K 7/24, C08J 3/20**

(21) Application number: **99109785.8**

(22) Date of filing: **18.05.1999**

(84) Designated Contracting States:
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE**
Designated Extension States:
AL LT LV MK RO SI

(72) Inventors:
• Dupire, Marc
7022 Mons (BE)
• Michel, Jacques
7181 Seneffe (BE)

(71) Applicant: **FINA RESEARCH S.A.**
B-7181 Seneffe (Feluy) (BE)

(54) Reinforced polymers

(57) Provided is a method for the production of a re-
inforced polymer, which method comprises:

(a) introducing carbon nanotubes into a polymer to

provide a mixture of the polymer and the nanotubes;
(b) stretching the mixture at or above the melting
temperature (T_m) of the polymer; and
(c) stretching the mixture in the solid state so as to
orient the carbon nanotubes.

EP 1 054 036 A1

Description

[0001] The present invention is concerned with a method for producing a reinforced polymer by introducing carbon nanotubes into the polymer. The invention also relates to reinforced polymers produced by the present methods and the use of carbon nanotubes for improving the mechanical properties of polymers.

[0002] It has been known for many years that blending fibres, such as carbon fibres, with polymers can significantly improve the mechanical properties of the blends (see Polymer Composites, April 1987, Vol. 8, No. 2, 74-81; J. Composite Materials, Vol. 3, October 1969, 732-734; and Polymer Engineering and Science, January 1971, Vol. 11, No. 1, 51-56). The advantage of carbon fibres is that they are very light, but despite this exhibit relatively great mechanical strength. In particular they exhibit very high stiffness.

[0003] More recently, since the discovery of Buckminsterfullerene (C_{60}), it has been found that carbon tubes (often termed carbon nanotubes because of their diminutive dimensions) having a structure related to the structure of C_{60} exist, which have the potential to be used in similar ways to carbon fibres. In particular, the structure of carbon nanotubes makes their aspect ratio (length/diameter, L/D) comparable to that of long fibres. Typically the aspect ratio of carbon nanotubes is > 10000 . Thus, the aspect ratio of carbon nanotubes is generally much greater than that of conventional short fibres, such as short glass fibres and short carbon fibres. In addition, the tubes can potentially be lighter than conventional carbon fibres, whilst being stronger and stiffer than the best conventional carbon fibres (see P. Calvert "Potential application of nanotubes" in Carbon Nanotubes, Editor T. W. Ebbeson, 297, CRC, Boca Raton, Florida 1997).

[0004] Depending on their diameter, helicity, and number of layers (single-wall v. multiple-wall) carbon nanotubes have electronic properties between those of conductors and semi-conductors. In addition, carbon nanotubes have great mechanical strength, being cited as having bending modulus values of from 1000-5000 GPa. Moreover they have been mentioned in connection with new, highly efficient, fracture micromechanisms which would prevent pure brittle failure with a concomitant low strain. Thus, carbon nanotubes have been envisaged for use in many applications in recent years (see P. Calvert "Potential application of nanotubes" in Carbon Nanotubes, Editor T. W. Ebbeson, 297, CRC, Boca Raton, Florida 1997; T. W. Ebbeson, "Carbon Nanotubes", Annu. Rev. Mater. Sci., 24, 235, 1994; Robert F. Service, "Super strong nanotubes show they are smart too", Science, 281, 940, 1998; and B. I. Yakobson and R. E. Smalley, "Une technologie pour le troisième millénaire: les nanotubes", La Recherche, 307, 50, 1998).

[0005] However, in the past when producing polyolefin composites by incorporating carbon nanotubes, tangling of the nanotubes and consequent randomising of the orientations of the nanotubes has caused problems (see M. S. P. Shaffer, X. Fan, A. H. Windle, "Dispersion of carbon nanotubes: polymeric analogies", poster 39, p. 317 in Proceedings of Polymer '98", September 1998, Brighton (UK); P. M. Ajayan, "Aligned carbon nanotubes in thin polymer films", Adv. Mater., 7, 489, 1995; H. D. Wagner, O. Lourie, Y. Feldman and R. Tenne, "Stress-induced fragmentation of multi-wall carbon nanotubes in a polymer matrix", Appl. Phys. Lett., 72 (2), 188, 1998; and K. Yase, N. Tanigaki, M. Kyotani, M. Yomura, K. Uchida, S. Oshima, Y. Kuriki and F. Ikazaki, Mat. Res. Soc. Symp. Proc., Vol. 359, 81, 1995). In particular, tangling can give rise to a reduction in the homogeneity of fibre/polymer blends since it is difficult for the fibres to distribute themselves evenly within the surrounding polymer matrix. This reduces the mechanical strength of the blends, since lack of homogeneity introduces weak points in a blend at positions where, for instance, there is a relatively low concentration of fibre and a high concentration of polymer. Moreover the randomising of the orientation of the fibres also reduces the mechanical strength of the blends. This is because (for example) the maximum resistance to strain in a given direction will be achieved when all of the fibres in the blend are oriented with their longitudinal axes aligned in that direction. The further that a blend deviates from such an ideal orientation, the less the resistance to strain of the blend in that direction. However, up to present it has not been possible to control the orientation of the fibres to a degree sufficient to improve mechanical properties.

[0006] It is an aim of the present invention to overcome the problems associated with the above blends and methods. Accordingly, the present invention provides a method for the production of a reinforced polymer, which method comprises:

- (a) introducing carbon nanotubes into a polymer to provide a mixture of the polymer and the nanotubes;
- (b) stretching the mixture at or above the melting temperature (T_m) of the polymer; and
- (c) stretching the mixture in the solid state so as to orient the carbon nanotubes.

[0007] The present invention further provides use of oriented carbon nanotubes in a polymer to reinforce the polymer.

[0008] In the present invention, the polymer is not particularly limited as long as the polymer can be oriented in the molten and in the solid state. In a preferred embodiment, the polymer is preferably a polyolefin, such as a polyethylene or a polypropylene or blends thereof. When the polyolefin is a polymer of an olefin monomer having 3 or more carbon atoms, such as polypropylene, the polyolefin may be isotactic or syndiotactic. A particularly preferred polymer is isotactic polypropylene, iPP. Other polymers which can be used in the present invention include polyesters, such as PETs and PEEKs, polyamides, PVCs, and polystyrenes.

[0009] The present invention is advantageous in that it succeeds in orienting the carbon nanotubes within the polymer such that their longitudinal axes are more aligned with each other than would otherwise be the case. In this invention 'orienting' is intended to mean a degree of disentangling of the carbon nanotubes and/or a degree of aligning of the carbon nanotubes. Not only are the nanotubes oriented, but also the individual polymer molecules undergo a degree of orientation in the present method. Orientation of the nanotubes leads to a greater homogeneity and less tangling in the resulting blends, and a consequent significant improvement in the mechanical properties of the blends. In particular, superior tensile modulus (as measured at 10 % strain, hereafter termed modulus (10 %)) and tenacity can be achieved by the present blends as compared with known blends, whilst still retaining a relatively high toughness (the product of tenacity and strain).

[0010] For these reasons, the reinforced polymers of the present invention are useful in a wide variety of applications involving the reinforcement of polymers, including use in fishing gear, tyres, safety belts, sewing thread, protective clothing, durable man-made fibre, and in cement paste, mortar or concrete. The reinforced polymers of the present invention are particularly useful in applications where high tenacity polyolefin fibres and filaments are required (see, for example, M. Ahmed, "Polypropylene Fibres - Science and Technology", Textile Science and Technology 5, High tenacity industrial yarns 389-403 and 665-681, Elsevier Amsterdam 1982).

[0011] The stretching procedure of the present method comprises two sequential steps: stretching the polymer/nanotube mixture in the molten state (step b) and subsequently stretching the solidified material (step c).

[0012] Stretching step (b) may comprise one of many traditional ways of processing polyolefins, including fibre melt spinning, extrusion, thermoforming and injection moulding. In a preferred embodiment of the present method, stretching step (b) comprises extruding the polymer/nanotube mixture. The extrusion step is carried out at a temperature at or above the melt temperature (T_m) of the polymer, so that the polymer is molten. Extrusion can be adapted to form a polymer fibre (e.g. by melt spinning) or to form a polymer sheet (e.g. by extrusion through a slit die or by compression moulding). In a preferred embodiment, the extrusion step is carried out by extruding the mixture through a spinneret to produce a fibre.

[0013] The method of producing the precursor composite (the mixture produced prior to solid state stretching) may be important, since it can affect the efficiency and the ease of orientation of the composites. For example, fast cooling isotactic polypropylene melt will produce the smectic crystalline form, which can deform much more easily in the solid state than the monoclinic crystalline form. Thus, in a preferred embodiment, after stretching in the molten state a desired cooling protocol is selected in order that the most desirable precursor composite is obtained.

[0014] In step (c), the polymer is preferably stretched at a temperature at or above its glass transition temperature, T_g , and below its melting temperature, T_m . Preferably this stretching step comprises solid state drawing. This typically involves drawing the polymer in the form of a fibre or sheet across a series of rollers. If desired the stretching step can be repeated one or more times. Optionally in the case of a polymer film, stretching can be performed biaxially, with longitudinal and transverse stretching taking place either sequentially or simultaneously. The stretching conditions can be chosen depending upon the nature of the polymer used, including the thermal characteristics, the molecular weight and/or the molecular weight distribution of the polymer. The conditions to be chosen for a specific polymer are readily determined according to standard solid state drawing procedures.

[0015] As mentioned above, carbon nanotubes are used for reinforcement in the present invention. By carbon nanotubes it is meant carbon tubes having a structure related to the structure of Buckminsterfullerene (C_{60}). Although often termed carbon nanotubes because of their diminutive dimensions, the carbon nanotubes used in the present invention need not necessarily have dimensions of the order of nanometers in size. The dimensions of the nanotubes can be much greater than this. However, it is preferred that the nanotubes are of a diameter from 1-50 nm, more preferably about 10 nm. Preferably the nanotubes are 1 μ m or more in length, more preferably about 10 μ m in length. Thus, it is preferred that the nanotubes are endowed with a high aspect ratio, having a length/diameter (L/D) of 100 or more, preferably 10^3 or more and most preferably 10^4 or more. Therefore, composites containing these nanotubes should, when the nanotubes are properly aligned, have mechanical properties which behave similarly to those of composites containing continuous carbon fibres.

[0016] Increasing the aspect ratio of the nanotubes (at constant nanotube volume fraction and orientation) leads to enhanced strength and stiffness in the composite. A long aspect ratio makes it less important to functionalise the polymeric matrix as well as the nanotubes to give good covalent binding and ensure good interfacial shear strength.

[0017] It should be noted that the use in the present invention of effectively non-continuous nanotubes (short in comparison to regular carbon fibres) rather than continuous fibres, allows access to typical processing techniques useful for thermoplastics. These techniques permit high throughput production and fabrication of high quality, complex shaped composites. Furthermore, due to their high strength and small diameter, the high aspect ratio of the nanotubes will not be decreased by typical polymer processing operations such as extrusion and injection. Thus polymer composites comprising nanotubes can provide the best of both worlds: high mechanical strength and ease of processing.

[0018] The quantity of carbon nanotubes added to a given quantity of polymer is not particularly limited. Typically less than 50 % wt. of carbon nanotubes or less is added to the polymer. Preferably 30 % wt. or less and more preferably

20 % wt. or less of nanotubes is added. It is most preferred that 5 % wt. or less of nanotubes is added. A very small quantity of nanotubes is capable of beneficially affecting the properties of a polymer, such that very small quantities can be used, depending on the intended use of the polymer. However, for most applications it is preferred that 0.1 % wt. of nanotubes or greater is added.

[0019] The method of manufacture of the polymers used in the present invention is not especially limited. In one preferred embodiment, the carbon nanotubes are mixed with the monomer, prior to carrying out the polymerisation procedure. Thus, in this embodiment the polymer is formed *in situ* in the presence of the carbon nanotubes.

[0020] The present invention extends to a reinforced polymer obtainable according to the methods of the present invention. The reinforced polymers of the present invention have superior mechanical properties, as has been discussed above. These properties are particularly marked in the case where the polymers are spun into fibres. Typically, fibres formed from the present reinforced polymers have a high modulus (10%), a high tenacity whilst still retaining a relatively high toughness. It is preferred that the modulus (10%), the tenacity and/or the toughness of fibres formed from the present reinforced polyolefin are greater by at least 15 %, as compared with the equivalent properties of the same polymer not comprising carbon nanotubes after undergoing the same stretching procedure.

[0021] Any additives typically introduced into polymers can be included in the present reinforced polymers, provided that the additives do not prevent the enhanced mechanical properties of the present polymers being obtained. Thus, additives such as pigments, anti-oxidants, UV-protective HALS, lubricants, anti-acid compounds, peroxides, grafting agents and nucleating agents can be included.

[0022] Embodiments of the present invention will now be described in the following by way of example only.

Examples

Synthesis of nanotubes

[0023] The nanotubes used in the present invention were obtained by catalytic decomposition of acetylene on a metallic catalyst (cobalt salt deposited onto zeolites or silica gels) according to K. Hernadi *et al*, Catalysis Letters, 48, 229, 1997. Silica-supported catalyst samples (prepared from Co-acetate solution of different pH) were tested in the decomposition of acetylene. Catalysts with various pre-treatments (different calcination temperature, time and atmosphere) were compared for nanotube yield and quality (quantity of turbostratic straight and helical tubes).

Apparatus

[0024] In the following examples, melt compounding of the polyethylene (PE) and polypropylene (PP) with the nanotubes was carried out using a MiniMax mixer (see B. Maxwell, "SPE J.", 28, 24, 1972).

[0025] Films were produced by compression moulding between Mylar® (Du Pont) PET sheets above the melting points of the polyolefins (typically below 200°C). Injection moulded tensile bars were produced with the MiniMax moulder.

[0026] Fibres were produced by blending and melting the polyolefin in the MiniMax, cutting pieces from an extruded strand, feeding the pieces into the barrel of a capillary rheometer (a CEAST 1000) and melting. The extruded strand was wound onto a rotating wheel, whose rotation speed determined the final diameter of the monofilament.

[0027] Solid state stretching was carried out on a MiniMat from Rheometrics Scientific, formerly produced by Polymer Laboratories, which is a miniature tensile tester equipped with an oven (see F. Biddlestone *et al* "The application of the MiniMat to plastic fracture: a review", Polymer Testing, 6, 163, 1986).

[0028] The titer of the fibre is the mass of the fibre per unit length, measured in g/10 km. The titer was determined using a Lenzing vibroscope.

Tensile Properties Measurements

[0029] The tensile properties of the melt-spun filament, the films, and the injection-moulded bars, were tested at room temperature in a tensile machine (a MiniMat for the films and the bars and a Lenzing for the fibres). The methods employed were carried out according to ISO 5079.

[0030] The conditions of testing were thus as follows:

Temperature 23°C

Gauge length 10 mm

Cross head speed 10 mm/min

Tensile Parameter Definitions

[0031] Rigidity was estimated at 10 % strain (10xforce at 10 % strain divided by the titer of the unstretched fibre). Force is expressed in cN.

5 [0032] The tenacity is the force at break divided by the titer of the fibre. For iPP (used in the following Examples) 1cN/Text is equivalent to 9 MPa.

[0033] The strain at break was expressed in %: $(L/L_0) \cdot 100$, with L_0 being the gauge length (10 mm). The maximum strain was 600 % and was limited by the machine maximum traverse length.

10 *Birefringence*

[0034] The birefringence (Δn) is defined as the difference between the refractive index (n) of a fibre measured parallel to its axis, and that measured perpendicularly to its axis. It was measured in the following Examples using a Berek compensator (Zeiss - Germany) and utilised according to the method described in "Polarised light microscopy: principles, instruments, applications", W. J. Patzelt, 3rd edition, E. Leitz Wetzlar (1985) p65-67.

15 [0035] Birefringence is a measure of degree of orientation in the polymer. The greater the birefringence, the greater the degree of orientation in the polymer.

20 *Stretch Ratio (SR)*

[0036] The solid state drawing carried out in the following Examples was conducted by selecting a specific SR for stretching. The SR is the ratio of the lengths of the fibre after and before the drawing step. It is equivalent to the ratio of the cross section (or titre) after and before the drawing step. Thus SR 2.5 refers to a 2.5 fold increase in the fibre length as a result of stretching. SR max refers to the maximum stretching possible without breaking the fibre. In each case in the following Examples the solid state drawing step was conducted at 110°C.

Example

30 [0037] To 10 g of isotactic polypropylene (iPP) reactor powder (having a melt flow index (MFI) of 10, determined at 230°C with a 2.16 kg load according to ISO 1133) 0.3 g (3 % wt. based on the weight of the polymer) of carbon nanotubes were added. Also added were 3000 ppm of anti-oxidant (B215 from Ciba) and 500 ppm of calcium stearate. The iPP powder, the additives and the nanotubes were well dispersed using a high shear mixer. 1.6 g of the blended material was introduced into the mixing chamber of a MiniMax mouder, which was heated at 220°C. The mixture was maintained at 220°C for 1 minute under an argon atmosphere and the axial rotor was rotated at 80 rpm for 2 minutes, allowing further intimate blending of the nanotubes and the iPP.

35 [0038] The above procedure was repeated for iPP in the absence of nanotubes and for iPP in the presence of 3 % wt. carbon black (blackpearl) in the place of nanotubes.

40 [0039] For each of the above three mixtures, a strand was extruded from the MiniMax and cut into small pieces. The pieces were fed into the heated cylinder of a CEAST 1000 capillary rheometer at a temperature of 185°C. The molten polymer was extruded through a capillary (length to diameter ratio of 5 to 1 mm) and the strand was wound onto a rotating wheel. For each of the above three mixtures, nominal 11 dTex fibres (approximately 40 µm in diameter) were produced.

[0040] Before subjecting the extruded fibres to a solid state drawing procedure, single fibres formed from each of the three mixtures were tested for their tensile properties and their titer (dTe=g/10 km of fibre). Titer was measured using a vibroscope. The results of the tests are shown in Table 1 under the heading Comparative Example.

45 [0041] A first orientation procedure in the solid semi-crystalline state was conducted on partially oriented melt-spun fibres in a MiniMat at 110°C with a slow strain rate (10 %/min with a gauge length of 10 mm). The maximum elongation was limited to a stretch ratio (SR) of 2.5. The fibres were easily stretched under these conditions (no breaking occurred during stretching). Single fibres formed from each of the three mixtures were again tested for their tensile properties. The results of the tests are shown in Table 1 under the heading Example A.

50 [0042] An alternative orientation procedure in the solid semi-crystalline state was conducted on unoriented melt-spun fibres at 110°C with a slow strain rate (10 %/min with a gauge length of 10 mm). The maximum elongation was limited to the maximum stretch ratio just before break (SR_{max}). Single fibres formed from each of the three mixtures were again tested for their tensile properties. The results of the tests are shown in Table 1 under the heading Example B.

55

Table 1

Example	Additive	Fibre fabrication	Ti (dTex)	SR	Δn (*1000)	e Break (%)	E 10 (cN/Tex)	E 5 (cN/Tex)	E 1 (cN/Tex)	Te (cN/Tex)
Comparative Example	None	melt-spun	11.3	1	10	600	4.7	---	---	11.4
	3 % C black	melt-spun	10.3	1	10.5	580	4.1	---	---	8.5
	3 % C nanotubes	melt-spun	6.3	1	11	500	4.2	---	---	9.4
Example A	None	SR 2.5, (110°C)	4.8	2.4	25	60	130	210	---	31.4
	3 % C black	SR 2.5, (110°C)	4.3	2.4	25	75	150	240	---	28
	3 % C nanotubes	SR 2.5, (110°C)	2.5	2.5	27	47	450	650	---	65
Example B	None	SR max, (110°C)	2.1	5.4	33	27	340	520	800	60
	3 % C black	SR max, (110°C)	2.1	4.9	33	27	340	520	800	61
	3 % C nanotubes	SR max, (110°C)	1	6.3	33	20	1570	1800	2300	268

Ti: Titer

SR: Stretch Ratio

 Δn : Birefringence

e Break: strain at break

E10: Secant modulus at 10 % strain

E5: Secant modulus at 5 % strain

E1: Secant modulus at 1 % strain

Te: tenacity at break (force at break/initial titer)

[0043] From Table 1, it can be seen that in the Comparative Example in which no solid state drawing was carried out, no improvement in tensile properties was seen and in fact a slight decrease in strain at break was observed.

[0044] However, performing moderate solid state stretching at an SR of 2.5 (Example A in Table 1) led to an improvement in tensile properties. A higher stiffness, tensile strength and energy-absorption capability was observed, whilst the strain at break remained within acceptable limits. Thus the E10 and E5 are improved by approximately a factor of 3 in comparison with fibres stretched under the same conditions which contain no additive or contain only carbon black.

[0045] When stretching the fibres to their maximum capability (SR max) (Example B in Table 1) an extremely high tenacity and E 10 are observed for the fibres containing nanotubes (approximately a 4-fold increase over the reference fibres). The strain at break remains within acceptable limits.

[0046] In the latter example, the tenacity is approximately twice the highest reported value for high tenacity iPP fibres (with a maximum tenacity of 130 cN/Text). The extension at break is still comparable to the reference fibres, hence the reinforced fibres are endowed with enhanced toughness. The stiffness could only be assessed at 1 % strain and was measured at approximately 2300 cN/Text (21 GPa). The theoretical upper modulus (based on 1.4 % volume fraction of perfectly aligned nanotubes in a highly oriented iPP fibre) should be between 38 GPa and 70 GPa depending on the value of the E modulus for the nanotubes (from 2000-5000 GPa).

[0047] Thus the reinforced polymers of the present invention have tensile properties comparable with the theoretical maximum obtainable values and are significantly improved over known reinforced polymers.

[0048] Furthermore, since excellent tensile properties are obtained for fibre comprising nanotubes which have been subjected to solid state drawing with an SR ratio of only 2.5, then thermal bonding in the normal processing temperature window is possible for iPP fibres of modest orientation. Thus, non-woven fabrics having high strength, stiffness and toughness can be produced using the present fibres.

Claims

1. A method for the production of a reinforced polymer, which method comprises:
 - (a) introducing carbon nanotubes into a polymer to provide a mixture of the polymer and the nanotubes;
 - (b) stretching the mixture at or above the melting temperature (T_m) of the polymer; and
 - (c) stretching the mixture in the solid state so as to orient the carbon nanotubes.
2. A method according to claim 1, wherein step (b) comprises extruding the mixture.
3. A method according to claim 2, wherein a polymer fibre or a polymer sheet is produced in the extrusion step.
4. A method according to any preceding claim, wherein step (c) comprises stretching the mixture at a temperature from the glass transition temperature (T_g) of the polymer to the melting temperature (T_m) of the polymer.
5. A method according to any preceding claim, wherein step (c) comprises solid state drawing.
6. A method according to any preceding claim, wherein 50 % by weight or less of carbon nanotubes is introduced into the polymer.
7. A method according to any preceding claim, wherein the carbon nanotubes are introduced into the polymer by contacting the nanotubes with a monomer and then polymerising the monomer in the presence of the nanotubes to form the mixture.
8. A method according to any preceding claim, wherein the polymer comprises a fibre-forming polymer
9. A method according to any preceding claim, wherein the polymer comprises a polyolefin, a polyester and/or a polyamide.
10. A method according to claim 9, wherein the polyolefin comprises a polyethylene and/or a polypropylene.
11. A reinforced polymer obtainable according to a method as defined in any preceding claim.
12. A reinforced polymer fibre or polymer she 1, formed from a reinforced polymer as defined in claim 11.

EP 1 054 036 A1

13. Use of a reinforced polymer fibre or polymer sheet as defined in claim 12 in fishing gear, a tyre, a safety belt, sewing thread, protective clothing, as a durable man-made fibre, or in cement paste mortar or concrete.
14. Use of oriented carbon nanotubes in a polymer to reinforce the polymer.
15. Use according to claim 14 in order to increase the tensile modulus and/or the tenacity and/or the toughness of the polymer.
16. Use according to claim 14 or claim 15, wherein the polymer comprises a polyolefin, a polyester and/or a polyamide.
17. Use according to claim 16, wherein the polyolefin comprises a polyethylene and/or a polypropylene.



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 99 10 9785

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
A	WO 97 15934 A (MITSUBISHI MATERIALS CORP ;HYPERION CATALYSIS INT (US); SHIBUTA DA) 1 May 1997 (1997-05-01) * page 5, line 38 - page 6, line 6 * * claim 1 *	1-17	C08K7/24 C08J3/20
A	GB 1 179 569 A (PIRELLI) 28 January 1970 (1970-01-28) * claims 1,3 *	1-17	
			TECHNICAL FIELDS SEARCHED (Int.Cl.7)
			C08K C08J
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 29 October 1999	Examiner Siemens, T
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	

EPO FORM 1503 03/82 (P/0401)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 99 10 9785

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

29-10-1999

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9715934 A	01-05-1997	JP 9111135 A	28-04-1997
		AU 7334696 A	15-05-1997
		EP 0857349 A	12-08-1998
GB 1179569 A	28-01-1970	US 3513067 A	19-05-1970

EPO FORM P4489

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82